











Prior Authorization Request Administrative Information

Member Information					
Last name	First name		МІ		
Member ID	Date of birth				
	X" or Intersex				
Current gender Female Male Transge	ender male 🔲 Tra	nsgender female Othe	-		
Place of residence Home Nursing facility	Other				
Race/ethnicity Preferred spoken la	anguage	Preferred written lang	uage		
MassHealth does not exclude people or treat them differently because of race, color, national origin, age, disability, religion, creed, sexual orientation, or sex (including gender identity and gender stereotyping).					
Plan Contact Information					
Please indicate the member's MassHealth Plan according to the Plan's contact information belo		his completed and signed	form		
MassHealth Fee-For-Service (FFS) Plan, Primary Care Clinician (PCC) Plan, Primary Care Accountable Care Organization (PCACO) Plan, Children's Medical Security Plan, and Health Safety Net Plan					
☐ MassHealth Drug Utilization Review Prog	gram				
Pharmacy: Fax: (877) 208-7428 - Tel: (800) 745-7318				
MassHealth Managed Care Organization	n (MCO) and Acco	untable Care Partnershi	p Plans (ACPP)		
☐ Fallon Health					
Online Prior Authorization: go.covermymeds.com/OptumRx					
Online Prior Authorization: providerportal.s	urescripts.net/Provi	derPortal/optum			
Pharmacy: Fax: (844) 403-1029 - Tel: (844) 720-0033					
☐ Health New England					
Online Prior Authorization: go.covermymeds.com/OptumRx					
Pharmacy: Fax: (800) 550-9246 - Tel: (800) 918-7545					
Online Prior Authorization (Non-Specialty Drugs): go.covermymeds.com/OptumRx					
Online Prior Authorization (Specialty/Medical Drugs): provider.massgeneralbrighamhealthplan.org					
Pharmacy: Fax: (844) 403-1029 - Tel: (800) 711-4555					
☐ Tufts Health Plan					
Online Prior Authorization: point32health.promptpa.com					
Pharmacy: Fax: (617) 673-0939 - Tel: (888) 257-1985					
☐ WellSense Health Plan					
Online Prior Authorization: wellsense.org/providers/ma/pharmacy/prior-authorizations					
Pharmacy: Fax: (833) 951-1680 - Tel: (877) 417-1822					

Inhaled Respiratory Agents Prior Authorization Request

MassHealth reviews requests for prior authorization (PA) on the basis of medical necessity only. If MassHealth approves the request, payment is still subject to all general conditions of MassHealth, including current member eligibility, other insurance, and program restrictions. MassHealth will notify the requesting provider and member of its decision. Keep a copy of this form for your records. If faxing this form, please use black ink.

Additional information about these agents, including PA requirements and preferred products, can be found within the MassHealth Drug List at **www.mass.gov/druglist**.

Medication information	
Medication requested (Check one or all that apply. W	here applicable, the brand name is provided in brackets
for reference.)	
Anticholinergics	Qvar Redihaler (beclomethasone inhaler)
☐ Lonhala (glycopyrrolate)	Long-acting Beta Agonists
☐ Yupelri (revefenacin)	arformoterol
Combination Products	☐ formoterol
☐ Airduo Digihaler (fluticasone/salmeterol)	Striverdi (olodaterol)
Airsupra (albuterol/budesonide)	Short-acting Beta Agonists
Bevespi (glycopyrrolate/formoterol)	albuterol inhaler ‡
☐ Breztri (budesonide/glycopyrrolate/formoterol)	levalbuterol inhalation solution
Duaklir (aclidinium/formoterol)	Proair Digihaler (albuterol inhalation powder)
fluticasone/salmeterol [Airduo Respiclick]	[‡] Brand name Proair, Proventil, and Ventolin are
Stiolto (tiotropium/olodaterol)	available without prior authorization.
☐ Trelegy (fluticasone furoate/umeclidinium/	*Brand name Flovent is available without prior
vilanterol)	authorization.
Corticosteroids	Other Medication
☐ Alvesco (ciclesonide inhaler)	
Armonair Digihaler (fluticasone propionate)	Other*
☐ Asmanex (mometasone) 110 mcg ≥ 12 years	*If request is for a non-preferred brand name or
Asmanex (mometasone) 220 mcg < 12 years	generic product, please attach supporting
☐ budesonide inhalation suspension ≥ 13 years	documentation (e.g., copies of medical records
☐ fluticasone propionate inhalation aerosol ≥ 12	and/or office notes regarding adverse reaction or
years [Flovent] ±	inadequate response to the preferred product).
fluticasone propionate inhalation powder	
[Flovent] ±	
Dose and frequency of medication requested	
Number of inhalers/month	
Indication (Check all that apply or include ICD-10 code	e, if applicable.)
Asthma (Specify severity below.)	
☐ Intermittent ☐ Mild Persistent	☐ Moderate Persistent ☐ Severe Persistent
☐ Chronic Obstructive Pulmonary Disease (COPD) (S	pecify severity and subtype below.)
Severity Mild Moderate Severe Ve	ery severe

PA-37 (Rev. 05/24) over

☐ Exercise-induced bronchospasm☐ Reactive airway disease	
Other	
Please list all other medications currently prescribed for the member for this indication.	
Is this member a referral candidate for care coordination? Yes No If yes, MassHealth will offer care coordination services to this member. Please describe which additional behavioral health services would be beneficial.	
Section I. Please complete for albuterol inhaler and Proair Digihaler requests.	_
1. For requests for albuterol inhaler, please attach medical records documenting an inadequate response or adverse reaction to an albuterol product available without prior authorization. *	
2. For requests for Proair Digihaler, has the member had a trial with brand name Proair, Proair Respiclick, Proventil, or Ventolin?	
Yes. Please list the dates/duration of trials, and outcomes in Section X.No. Please describe the clinical rationale why an albuterol inhaler is not appropriate for this member.	
* Brand name Proair, Proventil, and Ventolin do not require prior authorization.	_
Section II. Please complete for Asmanex 110 mcg requests in members ≥ 12 years of age and 220 mcg in members < 12 years of age.	1
Please describe the clinical rationale for the use of requested Asmanex strength in the requested age group.	
Section III. Please complete for all arformoterol, budesonide inhalation suspension, formoterol levalbuterol inhalation solution, Lonhala, and Yupelri requests.	٦l,
Please describe the medical necessity for a nebulized formulation.	7
2. For levalbuterol inhalation solution, has the member had a trial with albuterol solution?	
Yes. Please list the dates/duration of trials, and outcomes in Section X.	
No. Please describe the clinical rationale why albuterol solution is not appropriate for this member.	
3. For Lonhala and Yupelri, has the member had a trial with ipratropium inhalation nebulizer solution?	
 Yes. Please list the dates/duration of trials, and outcomes in Section X. No. Please describe the clinical rationale why ipratropium inhalation nebulizer solution is not appropria 	te
for this member.	
Section IV. Please complete for Airduo Digihaler, and fluticasone/salmeterol (generic Airduo Respiclick) requests.	_
 Has the member had a trial with fluticasone/salmeterol inhalation aerosol, powder (generic Advair)? Yes. Please list the dates/duration of trials and the outcomes in Section X. No. Please describe the clinical rationale for use of the requested agent in this member. 	
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 2. For Airduo Digihaler, has the member had a trial with fluticasone/salmeterol (generic Airduo Respiclick)' Yes. Please list the dates/duration of trials, and outcomes in Section X. No. Please describe the clinical rationale for use of the requested agent in this member. 	?			
Section V. Please complete for Alvesco, Armonair Digihaler, fluticasone propionate inhalati aerosol and powder (generic Flovent) [‡] , and Qvar Redihaler requests.	on			
Has the member had a trial with two inhaled corticosteroids?				
☐ Yes. Please list the dates/duration of trials, and outcomes in Section X.				
☐ No. Please document if there is a contraindication to all other inhaled corticosteroids.				
[‡] Brand name Flovent does not require prior authorization.				
Section VI. Please complete for Bevespi and Duaklir requests.				
Has the member had a trial with Anoro or Stiolto?				
☐ Yes. Please list the dates/duration of trials, and outcomes in Section X.				
No. Please describe the clinical rationale for use of the requested agent in this member.				
Section VII. Please complete for Trelegy requests.				
Has the member had a trial with fluticasone/vilanterol and Incruse or Anoro and Arnuity?				
Yes. Please list the dates/duration of trials, and outcomes in Section X.				
No. Please describe the clinical rationale for use of the requested agent in this member.				
Section VIII. Please complete for Breztri requests.				
Has the member had a trial with the following combination of the separate agents: Bevespi and Pulmicort inhalation powder?				
☐ Yes. Please list the dates/duration of trials, and outcomes in Section X.				
No. Please describe the clinical rationale for use of the requested agent in this member.				
Section IX. Please complete for Airsupra requests.				
Has the member had a trial with budesonide/formoterol or albuterol and Pulmicort inhalation powder?				
Yes. Please list the dates/duration of trials, and outcomes in Section X.				
☐ No. Please describe the clinical rationale for use of the requested agent in this member.				
Section X. Please complete as instructed in sections above.*				
Drug name Dates/duration of use				
Did the member experience any of the following? Adverse reaction Inadequate response Other				
Briefly describe details of adverse reaction, inadequate response, or other.				

Did	Dates/duration of use the member experience any of the following? Adverse reaction Inadequate response Other efly describe details of adverse reaction, inadequate response, or other.
	Dates/duration of use the member experience any of the following? Adverse reaction Inadequate response Other
	efly describe details of adverse reaction, inadequate response, or other. lease attach a letter documenting additional trials as necessary.
	ion XI. Please complete and provide documentation for exceptions to Step Therapy. Is the alternative drug required under the step therapy protocol contraindicated, or will likely cause an adverse reaction in, or physical or mental harm to the member? Yes No If yes, briefly describe details of contraindication, adverse reaction, or harm.
2.	Is the alternative drug required under the step therapy protocol expected to be ineffective based on the known clinical characteristics of the member and the known characteristics of the alternative drug regimen? Yes No If yes, briefly describe details of known clinical characteristics of member and alternative drug regimen.
3.	Has the member previously tried the alternative drug required under the step therapy protocol, or another alternative drug in the same pharmacologic class or with the same mechanism of action, and such alternative drug was discontinued due to lack of efficacy or effectiveness, diminished effect, or an adverse event? Yes No If yes, please provide details for the previous trial.
	Drug name Dates/duration of use Did the member experience any of the following? Adverse reaction Inadequate response Briefly describe details of adverse reaction or inadequate response.
4.	Is the member stable on the requested prescription drug prescribed by the health care provider, and switching drugs will likely cause an adverse reaction in or physical or mental harm to the member? Yes. Please provide details.
	□ No

Please continue to next page and complete Prescriber and Provider Information section.

Prior Authorization Request Prescriber and Provider Information

Prescriber Information		
Last name*	First name*	MI
NPI*	Individual MH Provide	er ID
DEA No.	Office Contact Name	
Address	City	State Zip
Email address		
Telephone No.*	Fax No.*	
* Required		
Please also complete for professionally	administered medication	ns, if applicable.
Start date	End date	
Servicing prescriber/facility name		☐ Same as prescribing provider
Servicing provider/facility address		
Servicing provider NPI/tax ID No.		
Name of billing provider		
Billing provider NPI No.		
Is this a request for recertification? Yes] No	
CPT code No. of visits	J code	No. of units
Prescribing provider's attestation, signal certify under the pains and penalties of perjoinformation section of this form. Any attached I certify that the medical necessity information complete, to the best of my knowledge. I under prosecution for any falsification, omission, or	ury that I am the prescribing I statement on my letterhead (per 130 CMR 450.204) on erstand that I may be subject concealment of any material	has been reviewed and signed by me. this form is true, accurate, and to civil penalties or criminal I fact contained herein.
Prescribing provider's signature		_
Printed name of prescribing provider (The form can either be signed by hand and		

(The form can either be signed by hand and then scanned, or it can be signed electronically using DocuSign or Adobe Sign. For electronic signatures, the signer can upload a picture of their wet signature. The typed text of a signature is not an acceptable form of an electronic signature.)